



DEPARTMENT OF HEALTH AND HUMAN SERVICES Public Health Service

Centers for Disease Control and Prevention INTER/INTRA-AGENCY AGREEMENT (IAA)



Payable Agreements (CDC is Procuring Agency)

1.	CDC IAA #: (10 to 13 digits)	2. PARTICIPAT	ING AGENCY IAA	#:	New Modificat	MENI ion Administrative		
	00FED05404-25	CPSC-IAG	G-01-1163 で	·Ś	Modification Number: 2			
4.	TITLE OF PROJECT:							
	National Electronic Injury Surveilland	e System (NEIS	S) Usefulness	of the NEIS	S-AIP to Produce A	nnual Stable Estima		
5.	DESCRIPTION OF WORK: (Please attach)		6. AMOU	JNT: (Not to exc	eed without written modif	ication)		
	See Attached \$ 165,000.00							
7.	NAME AND ADDRESS OF PARTICIPATING FEDERAL AGENCY:			LIAISON NAME	: :	PHONE #:		
	U.S. Consumer Product Safety Commission			Tom Schroe	der	(301) 504-7431		
	4330 East West Highway			EMAIL ADDRES	SS:	FAX #:		
	Bethesda, Maryland 20814-3724			TSchroeder(@cpsc.gov	(301) 504-0038		
8.	NAME AND ADDRESS OF CDC, CENTER, INSTITUTE OR OFFICE:			JAISON NAME	:	PHONE #:		
	National Center for Injury Prevention	• •		Chester L. P	ogostin	(770) 488-4805		
	4770 Buford Highway NE, MS-K59		. I	EMAIL ADDRES	SS:	FAX #:		
	Atlanta, Georgia 30341-3724		I.	clp3@cdc.go	ov	(770) 488-1665		
9.	9. PROJECT PERIOD: FUNDING PERIOD:							
			from: 10/0		through: 09/30	7/2006		
	: 10/01/2005 through: 09/30	1/2006	, from: 1070	71/2003	through: 09/30	3/2000		
	CDC AUTHORITY:							
	Economy Act approved June 30, 1932, as amended Other (Please specify) CDC DUNS # 9276	by 31 U.S.C. 1535 and 45465 CPSC Γ	l 1536 (See also ite NUNS# 069287	m #14) 7522				
		10.00 0.002						
11. PARTICIPATING AGENCY AUTHORITY:								
12.	CDC FUNDING INFORMATION: FOR CDC US	SE ONLY (CDC inter	nal form 6012 - mo	dified Document	History Racord)	ROPRIATION NUMBER:		
	T.C. FY DOC. REF.	DOC. NO.	CAN	O.C.	BACS	\$ AMOUNT		
Ad		(Original 10 digits) (Required)		(5 digits) (Required)	(10 digits) (For Budget			
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	050 05 214	00FED05404	6921 2306	25.3R	5609-21-11-01	\$165,000.00		
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6012 ADMINISTRATIVE APPROVAL NAME and EMAIL ADDRESS: (Please print) FMO BUDGET ANALYST SIGNATURE:								
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Health Service

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Payable Agreements (CDC is Procuring Agency)

CDC IAA #: 00FED05404-25

13. ADMINISTRATIVE BILLING REQUIREMENTS: CDC's ALC is 75090421. Other Agency's ALC: (required) 61000001

Billing is to be made through the use of the Online Payment and Collection (OPAC) system. Please include CDC's Official IAA # from Block #1 on all OPAC billings and correspondence. When CDC provides funds to the performing agency, in advance of receiving the

goods or services, the performing agency agrees to provide, within 15 day made during the quarter. The statements shall be provided to the following Desk, MS D-06, 1600 Clifton Road, Atlanta, GA 30333. (If recommended to the statement of the comment of the comm	ng address: DHHS, CDC, FMC), AP, Attn: ADVANCES/OPAC					
14. ADDITIONAL BILLING REQUIREMENTS: (This block must be completed if procuring services under the Economy Act.)							
All funds provided by CDC under this agreement must be obligated by the performing agency by the end of the FY in which the funds expire. Any unobligated but expired funds may not be used to fund services in subsequent periods. The CDC Financial Management Office (FMO) must be notified of any unobligated funds pertaining to this agreement at least 15 days before the end of the FY so that the agreement may be modified to reduce the funding amount when appropriate. This notification shall be provided to the following address: DHHS, CDC, FMO, AP, Attn: OPAC Desk, MS D-06, 1600 Clifton Road, Atlanta, GA 30333.							
15. PARTICIPATING AGENCY FUNDING and/or INFORMATION:							
(Please include name, telephone number, and email address of contact person.) Name: Email:							
	•	•					
Linda Murr (30)1) 504-0029	lmurr@cpsc.gov					
The participating agency as a signatory to the Common Rule states that in accepting these Interagency Agreement funds, it will abide by the human subjects research requirements stated in the Common Rule, and certify that all necessary assurances and institutional review board (IRB) approvals are obtained. The participating agency is NOT a signatory to the Common Rule. Upon issuance of these Interagency Agreement funds, it is the responsibility of the CDC Center, Institute, or Office (CIO) to certify that all necessary assurances and institutional review board (IRB) approvals are obtained. The CIO Associate Director for Science (ADS) must determine the Applicability of Human Subjects Regulations.							
17. OTHER REQUIREMENTS:							
A. Travel under this agreement is subject to allowances authorized in accordance with Federal Travel Regulations, Joint Federal Travel Regulations, and/or Foreign Service Regulations.							
B. CDC will retain the title to any equipment procured under this agreement, unless otherwise justified in the statement of work.							
18. CDC ACCEPTANCE: (please print)	19. PARTICIPATING AGENCY	ACCEPTANCE: (please print)					
Name: Ileana Arias, Ph.D.	Name: Donna Hutton						
Title: Acting Director, NCIPC	Title: Contracting Officer						
Email address: JA rig s@cdc.gov	Email address: dhutton@cpsc	.gov					
Signature:Sfellow for Date: 11/29/05	Signature:	Date: 12/23/15					
This agreement may be terminated by either agency upon a 30-day advance written notice. This agreement may be modified by mutual written consent of all parties.							

NTERAGENCY AGREEMENT BETWEEN THE CONSUMER PRODUCT SAFETY COMMISSION (CPSC) AND

THE CENTERS FOR DISEASE CONTROL AND PREVENTION (CDC) (00FED05404-25)

This document sets forth the terms of agreement for services, supplies, and/or material between the U.S. Consumer Product Safety Commission (CPSC) and the Department of Health and Human Services (DHHS), Centers for Disease Control and Prevention (CDC).

This document serves as an addendum to the Interagency Agreement (number 00FED05404) between the Centers for Disease Control and Prevention and the U.S. Consumer Product Safety Commission covering the expansion of the National Electronic Injury Surveillance System (NEISS) to collect data on all injuries. This system is called the NEISS All Injury Program (NEISS-AIP).

This addendum covers a special study entitled "Evaluating the Usefulness of the NEISS-AIP to Produce Annual Stable Estimates of the Incidence of Traumatic Brain Injury Treated in Emergency Departments in the United States". This project is outlined below. Additional documentation is attached as Appendices.

I. DESCRIPTION OF SERVICES

Background: A previously CDC-funded preliminary traumatic brain injury (TBI) surveillance pilot study investigated the feasibility of obtaining national estimates of the incidence of TBI treated in US hospital EDs, using NEISS AIP data, and the suitability of using NEISS-AIP data for ongoing surveillance of TBI treated in U.S. emergency departments. Specifically, the CPSC, in collaboration with CDC, reviewed and abstract data from medical records at a stratified, convenience sample of 10 NEISS-AIP hospitals to determine the best approach to identifying and characterizing ED cases with TBI and to assess the potential of NEISS-AIP data to distinguish true TBIs from other (extracranial) head injuries. The results of the pilot study indicate that NEISS-AIP could indeed be used to provide accurate and timely TBI data. It also showed that the addition of a second screen for identifying TBI cases from the NEISS-AIP could enhance the completeness of case identification by as much as 20 percent. The aforementioned "second screen" was produced in 2003 (in attached protocol, Appendix 1, Subappendix 1)). This study is an extention of the preliminary TBI surveillance pilot study. It will further investigate the most efficient and cost-effective case identification criteria and methods for full implementation of a TBI special study using all NEISS-AIP hospitals. This effort will provide statistically valid national estimates of the incidence of persons with TBIs treated in US hospital emergency departments.

Purpose:

This project is aimed at further evaluating the usefulness of NEISS-AIP to produce annual stable estimates of the incidence of TBI treated in hospital emergency departments (EDs) in the U.S. This project will begin in FY 2006.

Methods:

We propose to test case criteria for identifying persons with TBI treated in a sample of NEISS-AIP hospitals. This pilot study has two phases. <u>Phase I</u> will be used to pre-test the approach, proposed case criteria, and coder training material. Based on the results of Phase I, adjustments may be necessary prior to the next phase. <u>Phase II</u> will be used to evaluate the case criteria and feasibility of full implementation of a TBI Special Study using all NEISS-AIP hospitals.

TBI case criteria for this pilot study will go beyond the usual methods used by NEISS-AIP hospital coders. The new criteria will hopefully identify most if not all TBI cases treated at the sample NEISS-AIP hospitals regardless of whether or not the TBI was the principal diagnosis as determined by the attending ER physician. NEISS-AIP hospital coders will be asked to carefully review medical records for signs, symptoms, acronyms, or words commonly used by health care providers to characterize TBI or head injury in the medical record. The NEISS-AIP hospital coder will then record some of these terms in a brief narrative along with other data on circumstances of the injury incident. When this narrative is entered into the *first screen* on the NEISS-AIP electronic data entry system, a TBI special study *second screen* will be triggered. The NEISS-AIP hospital coder will then complete requested information about the TBI or head injury on the *second screen* using information from the patient's medical record. Information on the *second screen* will be used to determine true TBI cases for surveillance purposes. For more information, see the *Methods Section* of attached protocol. CPSC and CDC will use:

- 1. A proposed TBI standard case inclusion criteria (in attached protocol, Appendix 1);
- 2. The final version of the "second screen" (in attached protocol, Appendix 1, Subappendix 1)

II. DURATION OF AGREEMENT

This agreement is approved for one year from the date of signature for both agencies. If the results indicate that NEISS is able to produce annual stable estimates of TBI treated in ED, we will extend the agreement to five years.

III. ESTIMATED COSTS

Medical record review and abstraction for the "second screen" cost estimates: It is estimated that each year, 40,000 (forty thousand) medical records of possible TBI cases would be reviewed and abstracted to complete the "second screen" for our study; it is expected that at least 30,000 (approximately 80%) will meet the TBI case inclusion criteria. Each record will be reviewed at a cost of approximately \$4.00 per record; for an approximate total cost of \$157,000.00 per year.

Estimated annual costs are \$165,000.00. This cost estimate is broken down into the following sub-categories:



IV. <u>FUNDING</u>

All funds provided by CDC in this agreement must be obligated by the performing agency by the end of the fiscal year in which the funds expire. Any unobligated but expired funds may not be used to fund services in subsequent periods. The CDC Financial Management Office (FMO) must be notified of any unobligated funds pertaining to this agreement at least 15 days before the end of the fiscal year so that the agreement can be amended to reduce the obligated amount when appropriate. The notification must be provided to the address cited below (in paragraph V).

V. ACCOUNTING AND BILLING INFORMATION

Agency Agency Symbol Appropriation CAN Object Class UFMS Budget Activity	From CDC 75090421 7560943 6921 2306 25.3R 5609-21-11-01	<u>To</u> CPSC 61-00-0001 06 PS EXFM 4310 11179 252e		
Amount EIN No DUNS	\$165,000 58-6051157 927645465	\$165,000 52-0978750		
		. 069287522		

When billing CDC through the OPAC system, CPSC will reference agreement number 00FED05404-25.

When funds are provided to the performing agency in advance of services being performed or goods being delivered, the performing agency is required to provide, within 15 days of the end of each quarter, statements of obligations and expenditures made during the quarter. These statements are also provided to the address below:

CDC, FMO Attn: OPAC Desk 1600 Clifton Road, MS D-06 Atlanta, GA 30333

VI. EQUIPMENT

There is no equipment to be covered under this agreement.

VII. TRAVEL

There is no travel involved in this agreement.

VIII. CONFLICT WITH EXISTING AGREEMENTS

There is no duplication or conflict with existing agreements, policy, or statute.

IX. PROGRAM CONTACTS

CDC: Victor G. Coronado, MD, MPH NCIPC, DIDOP (F41) 4770 Buford Highway, NE Atlanta, Georgia 30341-3714 (770) 488-1568

CPSC: Tom Schroeder CPSC 4330 East West Highway. Rm 604D Bethesda, MD 20814-4408 (301) 504-0539 x1246

X. <u>BUDGET CONTACTS</u>

CDC: Deborah Mathis NCIPC/OD (K62) 4770 Buford Highway, NE Atlanta, Georgia 30341-3724 (770) 488-4695

CPSC: Donna Hutton Contracting Officer, CPSC 4330 East West Highway, Rm 517 Bethesda, MD 20814-4408 (301) 504-7009

X. MODIFICATION AND CANCELLATION

This agreement may be modified by mutual consent of both parties or canceled upon 60 days advance written notice by either party.

XI. AUTHORITY

This agreement is entered into under Section 601 of the Economy Act, as amended (31 U.S.C. 1535) and the Consumer Product Safety Act.